

Attorney Docket No.:  
Inventors:  
Serial No.:  
Filing Date:  
Page 3

KUZ0030US.NP.  
Toshimitsu et al.  
Not yet assigned  
Herewith

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This listing of the claims will replace all prior versions  
and listings of claims in the application:

**Listing of the claims:**

Claim 1 (original): A transdermal preparation  
containing pergolide and/or a pharmaceutically acceptable  
salt thereof, wherein said preparation is capable of  
achieving a plasma AUC ratio of pergolide or the  
pharmaceutically acceptable salt thereof to at least one  
metabolite thereof of 1:0.5 to 1:5.

Claim 2 (original): The transdermal preparation  
according to claim 1, wherein the plasma AUC ratio of  
pergolide and/or a pharmaceutically acceptable salt thereof  
to at least one metabolite thereof is 1:0.5 to 1:3.5.

Claim 3 (original): The transdermal preparation  
according to claim 2, wherein the plasma AUC ratio of  
pergolide and/or a pharmaceutically acceptable salt thereof  
to at least one metabolite thereof is 1:0.5 to 1:2.

Attorney Docket No.: KUZ0030US.NP.  
Inventors: Toshimitsu et al.  
Serial No.: Not yet assigned  
Filing Date: Herewith  
Page 4

Claim 4 (currently amended): The transdermal preparation according to ~~any one of claims 1 to 3~~ claim 1, wherein the metabolite is one or more kinds comprising pergolide sulfoxide, pergolide sulfone, despropyl pergolide or despropyl pergolide sulfoxide.

Claim 5 (original): The transdermal preparation according to claim 4, wherein the metabolite is pergolide sulfoxide.

Claim 6 (currently amended): The transdermal preparation according to ~~any one of claims 1 to 5~~ claim 1, wherein the pharmaceutically acceptable salt is one or more kinds comprising hydrochloride, sulfate, mesylate, citrate, fumarate, tartarate, maleate or acetate.

Claim 7 (original): The transdermal preparation according to claim 6, wherein the pharmaceutically acceptable salt is mesylate.

Claim 8 (currently amended): The transdermal preparation according to ~~any one of claims 1 to 7~~ claim 1, wherein the ratio (A/B) of the maximum plasma level (A) of

Attorney Docket No.: KUZ0030US.NP.  
Inventors: Toshimitsu et al.  
Serial No.: Not yet assigned  
Filing Date: Herewith  
Page 5

pergolide and/or the pharmaceutically acceptable salt thereof to the plasma level (B) thereof in the next administration and/or the ratio (A'/B') of the maximum plasma level (A') of pergolide sulfoxide to the plasma level (B') of pergolide sulfoxide in the next administration is less than 2.

Claim 9 (currently amended): The transdermal preparation according to ~~any one of claims 1 to 8~~ claim 1, wherein (meth)acrylic acid copolymer is contained in an adhesive layer.

Claim 10 (original): The transdermal preparation according to claim 9, wherein the acrylic polymer except (meth)acrylic acid copolymer is further contained in an adhesive layer.

Claim 11 (original): A transdermal preparation containing pergolide and/or the pharmaceutically acceptable salt thereof, wherein the ratio (A/B) of the maximum plasma level (A) of pergolide and/or the pharmaceutically acceptable salt thereof to the plasma level (B) thereof in the next administration and/or the ratio (A'/B') of the

Attorney Docket No.: KUZ0030US.NP  
Inventors: Toshimitsu et al.  
Serial No.: Not yet assigned  
Filing Date: Herewith  
Page 6

maximum plasma level (A') of pergolide sulfoxide to the  
plasma level (B') of pergolide sulfoxide in the next  
administration is less than 2.

Claim 12 (currently amended): The transdermal  
preparation according to ~~any one of claims 1 to 11~~ claim 1,  
wherein said preparation is an adhesive patch.